

This Page Is Inserted by IFW Operations  
and is not a part of the Official Record

## **BEST AVAILABLE IMAGES**

Defective images within this document are accurate representations of the original documents submitted by the applicant.

Defects in the images may include (but are not limited to):

- BLACK BORDERS
- TEXT CUT OFF AT TOP, BOTTOM OR SIDES
- FADED TEXT
- ILLEGIBLE TEXT
- SKEWED/SLANTED IMAGES
- COLORED PHOTOS
- BLACK OR VERY BLACK AND WHITE DARK PHOTOS
- GRAY SCALE DOCUMENTS

**IMAGES ARE BEST AVAILABLE COPY.**

**As rescanning documents *will not* correct images,  
please do not report the images to the  
Image Problem Mailbox.**

IFW

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

In Re the Application of:

STAMP

Serial No.: 10/767,860

Filed: January 28, 2004

Atty. File No.: 7095MH-2

For: "INJECTION DEVICE"

Commissioner for Patents  
P.O. Box 1450  
Alexandria, VA 22313-1450



SUBMISSION OF  
PRIORITY DOCUMENT

|   |
|---|
| <p>CERTIFICATE OF MAILING</p> <p>I HEREBY CERTIFY THAT THIS CORRESPONDENCE IS BEING DEPOSITED WITH THE UNITED STATES POSTAL SERVICE AS FIRST CLASS MAIL IN AN ENVELOPE ADDRESSED TO THE COMMISSIONER FOR PATENTS, P.O. BOX 1450, ALEXANDRIA, VA 22313-1450 ON <u>6/21/04</u></p> <p>BY: <u>Sheridan Ross P.C.</u></p> |
|---|

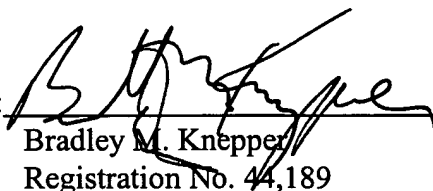
Dear Sir:

Enclosed is a certified copy of United Kingdom Patent Application No. 0401469.2, filed January 23, 2004, to support the previous claim of foreign priority benefits under 35 U.S.C. § 119 in connection with the above-identified application.

Respectfully submitted,

SHERIDAN ROSS P.C.

By:

  
Bradley M. Knepper  
Registration No. 44,189  
1560 Broadway, Suite 1200  
Denver, CO 80202-5141  
(303) 863-9700

Date: June 21, 2004



INVESTOR IN PEOPLE

The Patent Office  
Concept House  
Cardiff Road  
Newport  
South Wales  
NP10 8QQ

I, the undersigned, being an officer duly authorised in accordance with Section 74(1) and (4) of the Deregulation & Contracting Out Act 1994, to sign and issue certificates on behalf of the Comptroller-General, hereby certify that annexed hereto is a true copy of the documents as originally filed in connection with the patent application identified therein.

I also certify that the application is now proceeding in the name as identified herein.

In accordance with the Patents (Companies Re-registration) Rules 1982, if a company named in this certificate and any accompanying documents has re-registered under the Companies Act 1980 with the same name as that with which it was registered immediately before re-registration save for the substitution as, or inclusion as, the last part of the name of the words "public limited company" or their equivalents in Welsh, references to the name of the company in this certificate and any accompanying documents shall be treated as references to the name with which it is so re-registered.

In accordance with the rules, the words "public limited company" may be replaced by p.l.c., plc, P.L.C. or PLC.

Re-registration under the Companies Act does not constitute a new legal entity but merely subjects the company to certain additional company law rules.

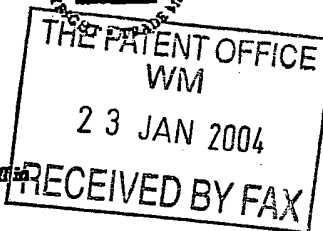
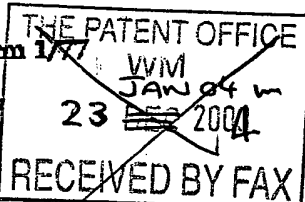
Signed

*Andrew Gersey*

Dated

4 February 2004

Patents Form 1/77

Patents Act 1977  
(Rule 16)23JAN04 E867832-2 D02973  
P01/7700 0,00-0401469.2 ACCOUNT CHA**Request for grant of a patent**

(See the notes on the back of this form. You can also get an explanatory leaflet from the Patent Office to help you fill in this form)

The Patent Office

Cardiff Road  
Newport  
South Wales  
NP10 8QQ

1. Your reference

P103497GB

2. Patent application number

(The Patent Office will fill this part in)

0401469.2

23 JAN 2004

3. Full name, address and postcode of the or of each applicant (underline all surnames)

The Medical House plc  
201 Newhall Road  
Attercliffe  
Sheffield  
S9 2QJ

Patents ADP number (if you know it)

8489239001

If the applicant is a corporate body, give the country/state of its incorporation

United Kingdom

4. Title of the invention

Injection Device

5. Name of your agent (if you have one)

Harrison Goddard Foote

"Address for service" in the United Kingdom to which all correspondence should be sent (including the postcode)

Fountain Precinct  
Balm Green  
SHEFFIELD  
S1 2JA

Patents ADP number (if you know it)

7914237001 /

6. Priority: Complete this section if you are declaring priority from one or more earlier patent applications, filed in the last 12 months.

Country

Priority application number  
(if you know it)Date of filing  
(day / month / year)

7. Divisionals, etc: Complete this section only if this application is a divisional application or resulted from an entitlement dispute (see note d)

Number of earlier UK application

Date of filing  
(day / month / year)

8. Is a Patents Form 7/77 (Statement of inventorship and of right to grant of a patent) required in support of this request?

Answer: YES if:

yes

- a) any applicant named in part 3 is not an inventor, or
- b) there is an inventor who is not named as an applicant, or
- c) any named applicant is a corporate body.

Otherwise answer NO (See note d)

Patents Form 1/77

## Patents Form 1/77

9. Accompanying documents: A patent application must include a description of the invention. Not counting duplicates, please enter the number of pages of each item accompanying this form:

Continuation sheets of this form

|             |    |
|-------------|----|
| Description | 12 |
| Claim(s)    | 3  |
| Abstract    | 1  |
| Drawing(s)  | 12 |

10. If you are also filing any of the following, state how many against each item.

Priority documents

Translations of priority documents

Statement of inventorship and right to grant of a patent (Patents Form 7/77)

Request for a preliminary examination and search (Patents Form 9/77) 1

Request for a substantive examination (Patents Form 10/77)

Any other documents (please specify)

11. I/We request the grant of a patent on the basis of this application.

Signature(s)

*Vanessa J Stainthorpe*

Date 23 January 2004

12. Name, daytime telephone number and e-mail address, if any, of person to contact in the United Kingdom

Vanessa J Stainthorpe

0114 274 3700

## Warning

After an application for a patent has been filed, the Comptroller of the Patent Office will consider whether publication or communication of the invention should be prohibited or restricted under Section 22 of the Patents Act 1977. You will be informed if it is necessary to prohibit or restrict your invention in this way. Furthermore, if you live in the United Kingdom, Section 23 of the Patents Act 1977 stops you from applying for a patent abroad without first getting written permission from the Patent Office unless an application has been filed at least 6 weeks beforehand in the United Kingdom for a patent for the same invention and either no direction prohibiting publication or communication has been given, or any such direction has been revoked.

## Notes

- If you need help to fill in this form or you have any questions, please contact the Patent Office on 08459 500505.
- Write your answers in capital letters using black ink or you may type them.
- If there is not enough space for all the relevant details on any part of this form, please continue on a separate sheet of paper and write "see continuation sheet" in the relevant part(s). Any continuation sheet should be attached to this form.
- If you have answered YES in part 8, a Patents Form 7/77 will need to be filed.
- Once you have filled in the form you must remember to sign and date it.
- Part 7 should only be completed when a divisional application is being made under section 15(4), or when an application is being made under section 8(3), 12(6) or 37(4) following an entitlement dispute. By completing part 7 you are requesting that this application takes the same filing date as an earlier UK application. If you want the new application to have the same priority date(s) as the earlier UK application, you should also complete part 6 with the priority details.

## INJECTION DEVICE

This invention relates to the field of injection devices for the administration of liquid medication, for example, insulin or growth hormone.

One type of injection device is known as a mini-needle or micro-needle device. These devices comprise a pressurised ("forced") injection system and have a needle which is shorter than that of conventional needle systems. The needle is normally hidden which is advantageous both for avoiding needle stick injuries and for minimising trauma to needle-phobic patients. The needle is hidden both before and after the injection is delivered, appearing only for the duration of the injection. Mini needle devices can typically deliver a larger volume of medication than needle-free devices and can deliver faster than conventional needle systems.

One such known device is described in WO00/09186 (Medi-Ject Corporation) for "Needle assisted jet injector" and this document gives a useful summary of prior art devices.

The device of WO 00/09186 includes a needle which is, in one embodiment, retractably located within an injector nozzle assembly. Upon activation of a force generating source, a portion of the needle extends past the nozzle assembly and penetrates the outer layer of skin to deliver medicament via jet injection to a deeper region. After activation, the needle retracts back into the nozzle assembly. The retractable needle is housed within the nozzle and is pushed forward so that it emerges in order to deliver an injection by the liquid medicament itself, when the medicament is itself pushed forward by

the plunger.

According to a first aspect of the present invention there is provided an injection device comprising an outer  
5 housing inside which is located

a barrel for holding a dose of a medicament;

a needle at one end the barrel and fixed with respect thereto, the needle and barrel being such that at least part of the needle is axially moveable in and out of said  
10 outer housing but is biased to be normally wholly inside said housing;

a plunger, axially moveable within the barrel;  
an inner housing intermediate the outer housing and the barrel and plunger; and

15 an energy source in communication with said inner housing,

wherein the inner housing is moveable by the energy source between three positions, namely

a first position in which the inner housing is in  
20 communication with both the plunger and the barrel such that, in use, the plunger and barrel are movable axially so as to move at least part of said needle out of the outer housing;

a second position in which the inner housing is in  
25 communication with the plunger but not the barrel such that, in use, said plunger is movable axially into said barrel so as to expel medicament through the needle; and

a third position in which the inner housing is in communication with neither the plunger nor the barrel  
30 such that, in use, the plunger and barrel are able to retract in order to retract the needle into the outer housing.

The injection device according to the present invention  
35 provides a simple and cost-effective means of delivering

medicament through a retractable mini needle. The device is able to deliver medicament to a depth beyond the length of the needle because of the propulsive force provided by the energy source.

5

The device requires that the needle (and hence also the barrel to which it is fixed) is moved axially so that the needle can appear beyond the end of the nozzle for the duration of the injection, after which the needle  
10 retracts automatically, out of sight of the user. The device also requires that the plunger is moved axially (into the barrel) so that medicament is ejected. The overall complexity of the injection device is significantly reduced by both of these requirements being  
15 effected by one component, namely the inner housing.

Preferably, said inner housing includes one or more flexible tags, biased radially inwardly by communication with said outer housing.

20

Preferably, one or more of said tags are situated at the rear end of the inner housing and are biased radially inwardly into communication with the plunger. Each rear tag may be moveable out of communication with the plunger  
25 when aligned with a corresponding recess in the outer housing. Preferably, each rear tag is substantially T-shaped. One leg of the T-shape enables the rear tag to hook over the plunger and, effectively, pull the plunger forward (in the first and second positions mentioned  
30 above). The other leg of the T-shape enables the rear tag to spring radially outwardly to catch in a recess in the housing (in the third position mentioned above).

Preferably, one or more of said tags are situated at the  
35 forward end of the inner housing and are biased radially



inwardly into communication with the barrel. Each forward tag may be moveable out of communication with the barrel when aligned with a corresponding recess in the outer housing. Preferably, each rear tag is substantially L-shaped.

In a preferred embodiment, said energy source is a compressed gas.

10 Preferably, the injection device further includes means for allowing the inner housing to move axially only forward with respect to the outer housing. Ideally, said means is an arrangement of serrations intermediate the housings.

15

Preferably, said needle is biased to be normally wholly inside said housing by means of a spring intermediate the barrel and the outer housing.

20 In one embodiment, said needle, barrel and plunger are removable from said device.

25 Preferably, the injection device further includes a removable needle cover which protects the needle during storage and before use. Advantageously, said needle cover includes means for pulling a protective rubber sheath or the like from said needle when said needle cover is removed from the device.

30 According to a second aspect of the invention, there is provided a method of delivering an injection using an injection device as described in any of the preceding paragraphs.

35 Preferred embodiments of the present invention will now

be more particularly described, by way of example only, with reference to the accompanying drawings wherein:

5 Figure 1 is a perspective view, partly in section, showing the injection device, in the condition in which it is supplied to a user, apart from the needle cover;

Figure 2, drawn to a larger scale, shows detail of part of the device shown in Figure 1;

10

Figure 3 is a perspective view, partly in section, showing the injection device, during an injection;

15 Figure 4, drawn to a larger scale, shows detail of part of the device shown in Figure 3;

Figure 5 is a perspective view, partly in section, showing the injection device, with the plunger fully depressed into the barrel;

20

Figure 6, drawn to a larger scale, shows detail of part of the device shown in Figure 5;

25 Figure 7 is a perspective view, partly in section, showing the injection device, after use and safe to dispose of;

Figure 8, drawn to a larger scale, shows detail of part of the device shown in Figure 7;

30

Figure 9 is a perspective view of the device, including the needle cover;

35 Figure 10 is perspective view, partly in section, showing detail of the needle cover;

Figure 11 is perspective view, partly in section, showing detail of the needle cover part way through being removed from the injection device; and

5

Figure 12 is a schematic view showing the relationship between tags 7A and ram 4, in one embodiment of the invention.

10 Throughout the following description, reference to a "forward" direction means the direction which is towards the patient when the injection device is in use. The "forward" end of the injection device is the end nearest the patient's skin when the device is in use. Similarly,  
15 reference to a "rearward" direction means the direction which is away from the patient and the "rearward" end of the device is the end furthest from the patient's skin when the injection device is in use.

20 Figure 1 is a perspective view, partly in section, showing the injection device, in the condition in which it is supplied to a user, apart from the needle cover (which is described below after the main operation of the device).

25

The principal components of the device will now be described with reference to Figures 1 and 2. An energy source 1 is provided at the rear of the device which, in this embodiment, is a gas cylinder similar to the type  
30 used in a conventional aerosol can or the like i.e. having a valve through which gas can be released at will and in a controlled manner. In an alternative embodiment of the invention, it is envisaged that a spring could be used as the energy source in place of a gas cylinder.

35

7

The valve 2 of the gas cylinder opens into a chamber 3, which in Figure 1 is of relatively small volume. The front wall of the chamber 3 is defined by a ram 4 which has an annular seal 5 at the rear thereof in order to make the chamber 3 gas-tight. The rear wall of the chamber 3 is defined by the back face of a generally cylindrical chamber housing 6.

The forward part of the ram 4 abuts or alternatively is integrally-formed with an inner housing 7 which closely surrounds a plunger 8 (and can therefore be referred to as the "plunger housing"). The rear of the plunger housing includes four orthogonally placed tags 7A, which each have a "hammer head" or T-shape and whose tendency to spring radially outwardly is restricted by the diameter of the chamber housing 6. If the ram 4 is integrally formed with the plunger housing 7 as illustrated in Figure 12, the tags 7A are positioned at the end of flexible legs cut into the housing, so that the tags 7A can move radially, with respect to the ram 4 and remainder of housing 7.

The hammer head of each tag 7A hooks over the enlarged head 8A of the plunger 8, so that the tags 7A are in contact with the plunger head 8A, as shown best in Figure 2.

The plunger 8 is the plunger of a syringe arrangement comprising a barrel 9 in which a predefined dose of liquid medicament is supplied and a relatively short needle 10 through which the medicament can be delivered to the patient. A nozzle 11 at the front end of the injection device normally conceals the needle 10 from the user's view. A spring 12, positioned between the outer housing and the barrel 9 biases the needle to be normally

wholly within the nozzle 11.

At the front end of the plunger housing 7, there are further orthogonally placed tags 7B, which each have a generally L shape and whose tendency to spring radially outwardly is restricted by the diameter of the chamber housing 6. The tags 7B each abut the flange at the rear of barrel 9.

There are four main stages in the operation of the device. Stage 1 is the condition shown in Figures 1 and 2, i.e. the device as supplied to a user, and as described above. The medicament is already present in the barrel 9 and the needle 10 is concealed from view within the nozzle 11. The plunger 8 is fully withdrawn from the barrel 9 (because of the liquid medicament contained within the barrel) and the head of the plunger 8A abuts the tags 7A. The rear of the remainder of housing 7 abuts the ram 4. The chamber 3 is of minimal volume.

Stage 2 of operation is the injection stage illustrated in Figures 3 and 4. With the injection device held against the patient's skin at the injection site, downward force is applied to the device in the direction indicated by the arrow F in Figure 2. This force causes the valve of the gas cylinder 1 to open, releasing gas into the chamber 3. As the chamber 3 fills with gas, the ram 4 is urged forward, consequently urging tags 7A against the plunger 8. As the tags 7A, and hence the plunger housing 7, are urged forward, the cooperation of the tags 7B against the barrel 9 means that the barrel is also urged forward, against the bias of a spring 12 (shown in Figure 3). As the barrel 9 moves forward, so does the needle 10 which is attached thereto and so the needle protrudes out of the nozzle 11 sufficiently to

9

enable an injection to be delivered. Therefore, initially, the ram 4 causes the plunger housing 7, the plunger 8, the barrel 9 and the needle 10 to move forwards.

5

Shortly after the plunger housing 7 starts to move forward, the tags 7B reach a lip in the chamber housing 6. The tags 7B spring radially outwardly over this lip, as shown in Figures 3 and 4. Once the tags 7B have sprung outwardly in this way, they are no longer in abutment with the barrel 9. This means that the barrel 9 (and hence needle 10) is no longer urged forwards because the forwardly-moving plunger housing 7, including tags 7B, are free to continue moving forward without contacting the barrel 9.

15

Therefore, once the device has reached the condition illustrated in Figures 3 and 4, continued forward movement of the ram 4 and plunger housing 7 causes the plunger 8 to be urged forward into the barrel 9. This expels the liquid medicament from the barrel 9, through the needle 10 to deliver an injection. It is the cooperation of the tags 7A with the enlarged head 8A of the plunger which transmits the forward force from the ram 4/housing 7 to the plunger 8.

25

The third stage in the operation of the device is illustrated in Figures 5 and 6. When the plunger 8 is fully depressed into the barrel 9, the full dose of medicament has been delivered into the patient. At this point, the tags 7A reach recesses 13 cut into the plunger housing 6 whereupon they are able to spring radially outwardly into those recesses (as illustrated in Figures 5 and 6). This outward movement of the tags 7A means that the "hammer head" shape is no longer in contact with

35

10

the enlarged head 8A of the plunger 8 and therefore the plunger 8 is no longer driven forward by the ram 4 and tags 7A. The plunger housing 7 may continue further forward until an end stop is reached.

5

The final stage in the operation of the device is illustrated in Figures 7 and 8. With the tags 7A located within recesses 13, neither the plunger 8 or the barrel 9 is impeded by any part of the plunger housing 7. Therefore the spring 12, which had been compressed by the forward motion of the barrel 9, urges the barrel 9 and hence the plunger 8 backwards until the ram 4 prevents further backward movement thereof. The backward movement is sufficient to cause the needle 10 to retract into the nozzle 11 so that it is no longer visible to the user and safe from the risk of causing a needle-stick injury. The used injection device can then be safely disposed of.

Blow-back is prevented by the provision of serrations 14 which guide the relative movement of the chamber housing 6 and the outermost housing. These serrations only permit relative movement in one direction, i.e. the chamber housing 6 moving forward with respect to the outermost housing.

25

Figure 9 is a perspective view of the injection device which, in this Figure, includes a needle cover 15. The needle cover is shown in further detail in Figures 10 and 11 and is used to protect the needle end of the injection device during transit, storage and before use to deliver an injection.

Regulations require that the needle (which is in direct communication with the medicament in the barrel) is sealed from the outside environment before use. This is

35

achieved by providing protection in the form of a rubber moulding 16 which covers the end of the needle, the rubber moulding 16 being surrounded by a nylon sheath 17. The rubber moulding and nylon sheath (the "needle protection") are fixed with respect to one another by a friction fit between one or more protrusions 16A on the rubber moulding and a corresponding one or more recesses 17A in the nylon sheath.

The nylon sheath and rubber moulding are so firmly fixed on the needle 10 that it is difficult, if not impossible, for a patient to pull them from the needle using his/her fingers alone. Therefore an outer needle cover 15 is provided which not only improves the aesthetic appearance of the injection device, before use, but also serves the function of facilitating the removal of the nylon sheath and rubber moulding.

The needle cover 15 is releasably retained on the front end of the injection device by the fit of annular protrusions 18 on part of the device housing with grooves 19 on the interior of the needle cover. The protrusions 18 and corresponding grooves 19 preferably extend around two equally opposed 60° portions of the circumference of the nozzle 11.

The grooves 19 are located on one or more (preferably equally spaced) flexible legs 20 which are flexible compared to the rest of the needle cover 15, about point P shown in Figure 9. Forward of each groove 19 is provided an inwardly projecting tab 21 on each flexible leg 20. Each tab 21 abuts the rear of the nylon sheath 17.

Turning now to Figure 11, when it is desired to remove



the needle cover 15 from the device, the user grips the needle cover, preferably in a region having texture or other grip-improving means 22, and pulls in the direction indicated by the arrow in Figure 11. The flexibility of the legs 20 permits the needle cover to ride over the protrusions 18, disengaging them from the grooves 19. The flexibility of the legs 20 is minimal enough not to cause the tabs 21 to become disengaged from the rear of the nylon sheath 17.

10

Therefore, as the needle cover 15 is pulled in the direction indicated by the arrow, the tabs 21 are urged against the rear of the nylon sheath 17 and sufficient force can be applied thereby to disengage the needle 10 from the rubber moulding 16. In this way, the entire moulding 16, nylon sheath 17 and needle cover 15 can be removed from the injection device and discarded, so that the injection device is then ready to use.

## CLAIMS

1. An injection device comprising an outer housing  
inside which is located  
5 a barrel for holding a dose of a medicament;  
a needle at one end the barrel and fixed with  
respect thereto, the needle and barrel being such  
that at least part of the needle is axially moveable  
in and out of said outer housing but is biased to be  
10 normally wholly inside said housing;  
a plunger, axially moveable within the barrel;  
an inner housing intermediate the outer housing and  
the barrel and plunger; and  
an energy source in communication with said  
15 inner housing,  
wherein the inner housing is moveable by the energy  
source between three positions, namely  
a first position in which the inner housing is  
in communication with both the plunger and the barrel  
20 such that, in use, the plunger and barrel are movable  
axially so as to move at least part of said needle out of  
the outer housing;  
a second position in which the inner housing is  
in communication with the plunger but not the barrel such  
25 that, in use, said plunger is movable axially into said  
barrel so as to expel medicament through the needle; and  
a third position in which the inner housing is  
in communication with neither the plunger nor the barrel  
such that, in use, the plunger and barrel are able to  
30 retract in order to retract the needle into the outer  
housing.
2. An injection device as claimed in claim 1 wherein  
said inner housing includes one or more flexible  
35 tags, biased radially inwardly by communication with

14

said outer housing.

- 5           3. An injection device as claimed in claim 2 wherein  
            one or more of said tags are situated at the rear  
            end of the inner housing and are biased radially  
            inwardly into communication with the plunger.
- 10           4. An injection device as claimed in claim 3 wherein  
            each rear tag is moveable out of communication with  
            the plunger when aligned with a corresponding recess  
            in the outer housing.
- 15           5. An injection device as claimed in claim 4 wherein  
            each rear tag is substantially T-shaped.
- 20           6. An injection device as claimed in claim 2 wherein  
            one or more of said tags are situated at the forward  
            end of the inner housing and are biased radially  
            inwardly into communication with the barrel.
- 25           7. An injection device as claimed in claim 6 wherein  
            each forward tag is moveable out of communication  
            with the barrel when aligned with a corresponding  
            recess in the outer housing.
- 30           8. An injection device as claimed in claim 7 wherein  
            each rear tag is substantially L-shaped.
- 35           9. An injection device as claimed in any of the  
            preceding claims wherein said energy source is a  
            compressed gas.
10. An injection device as claimed in any of the  
            preceding claims further including means for  
            allowing the inner housing to move axially only

15

forward with respect to the outer housing.

- 5           11. An injection device as claimed in claim 10 wherein  
said means is an arrangement of serrations  
intermediate the housings.
- 10           12. An injection device as claimed in any of the  
preceding claims wherein said needle is biased to be  
normally wholly inside said housing by means of a  
spring intermediate the barrel and the outer  
housing.
- 15           13. An injection device as claimed in any of the  
preceding claims wherein said needle, barrel and  
plunger are removable from said device.
- 20           14. An injection device as claimed in any of the  
preceding claims further including a removable  
needle cover which protects the needle during  
storage before use.
- 25           15. An injection device as claimed in claim 14 wherein  
said needle cover includes means for pulling a  
protective rubber sheath or the like from said  
needle when said needle cover is removed from the  
device.
- 30           16. An injection device substantially as described  
herein with reference to and as illustrated in any  
appropriate combination of the accompanying  
drawings.
- 35           17. A method of delivering an injection using an  
injection device as described in any of the  
preceding claims.

## ABSTRACT

An injection device comprising an outer housing inside which is located a medicament-holding barrel with a needle at one end thereof, at least part of the needle being moveable in and out of the outer housing, a plunger moveable within the barrel, an inner housing intermediate the outer housing and the barrel and plunger and an energy source in communication with said inner housing. The inner housing is moveable by the energy source between

a first position in which the plunger and barrel are movable axially so as to move at least part of said needle out of the outer housing;

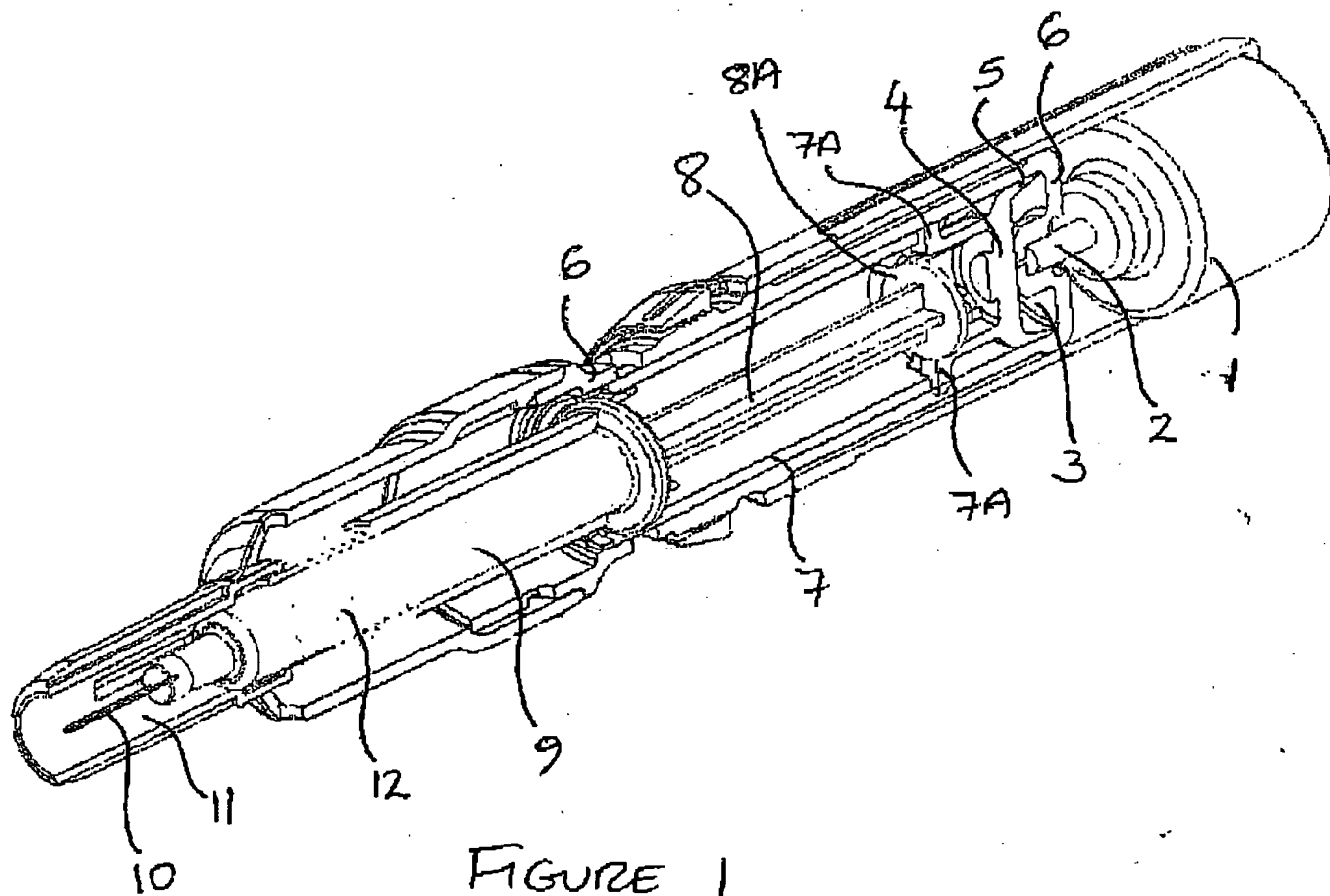
a second position in which the plunger is movable axially into said barrel so as to expel medicament through the needle; and

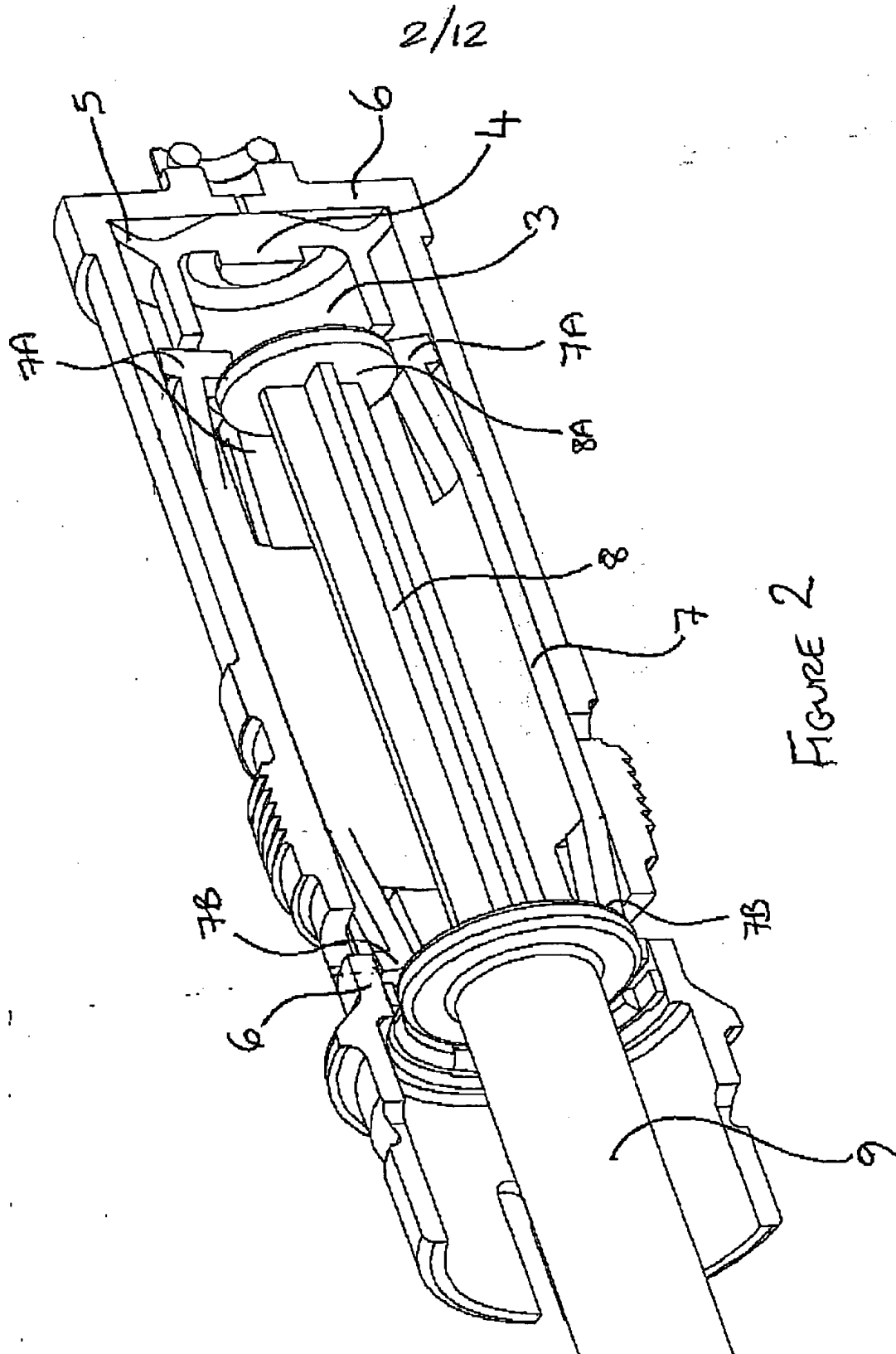
a third position in which the plunger and barrel are able to retract in order to retract the needle into the outer housing.

[Figure 1]

NOT FORWARDED

1/12





3/12

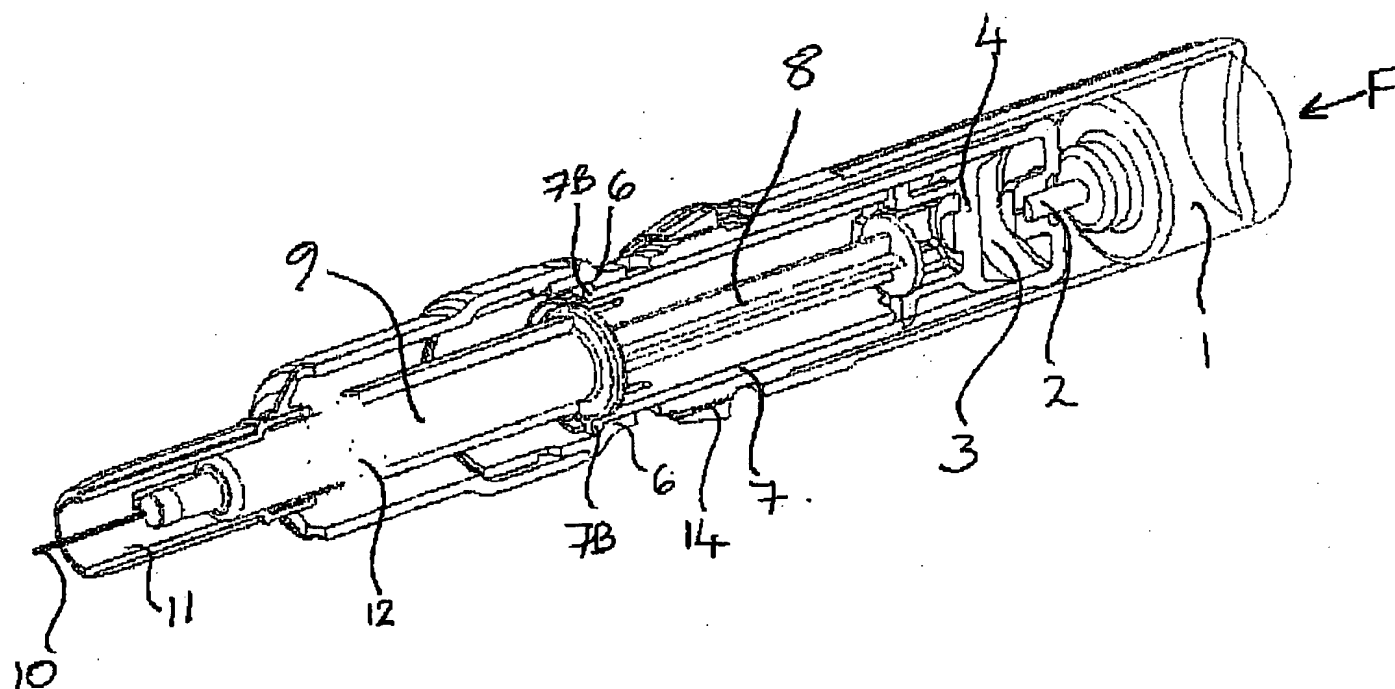


FIGURE 3



4/12

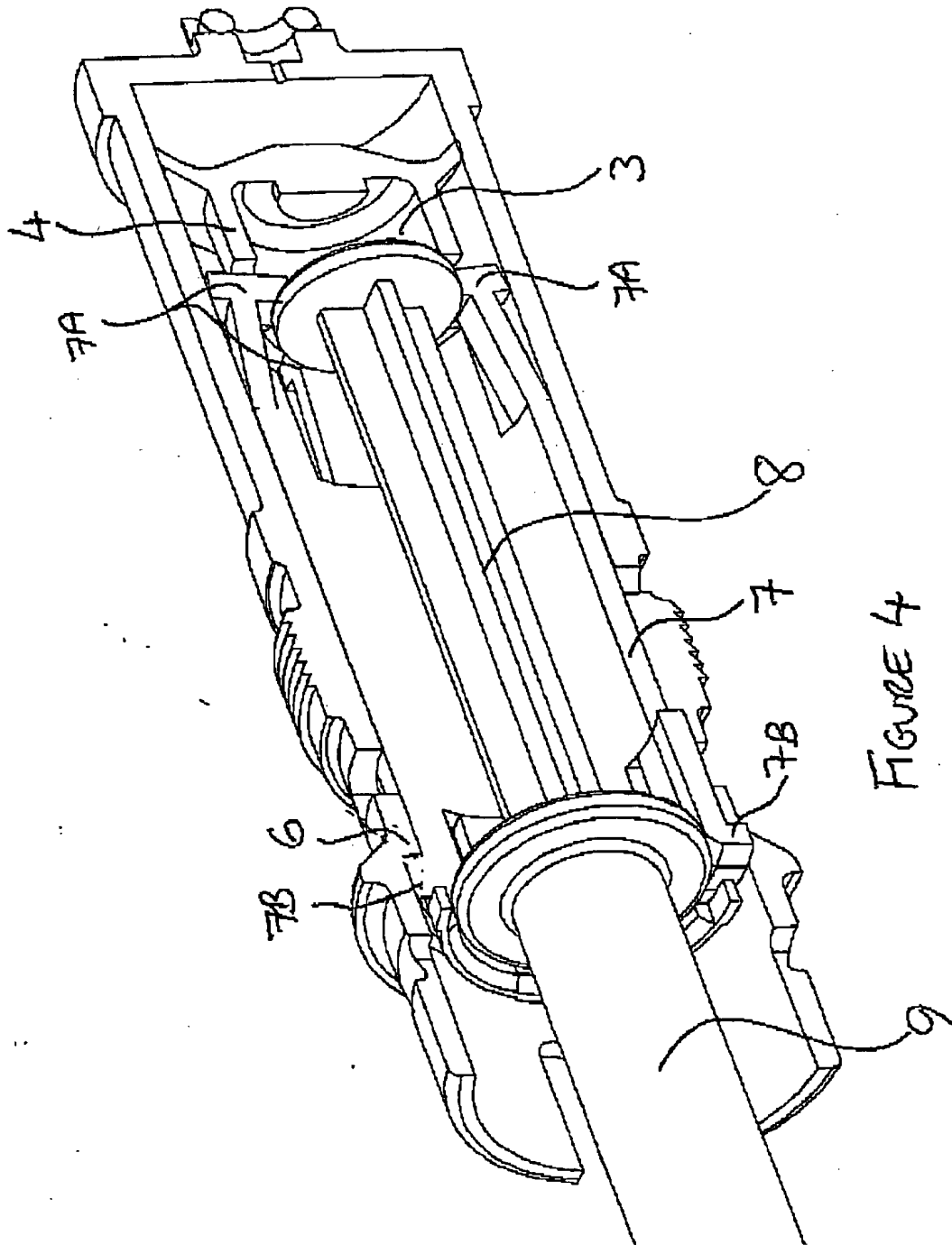


FIGURE 4

NOT TO BE REPRODUCED

5/12

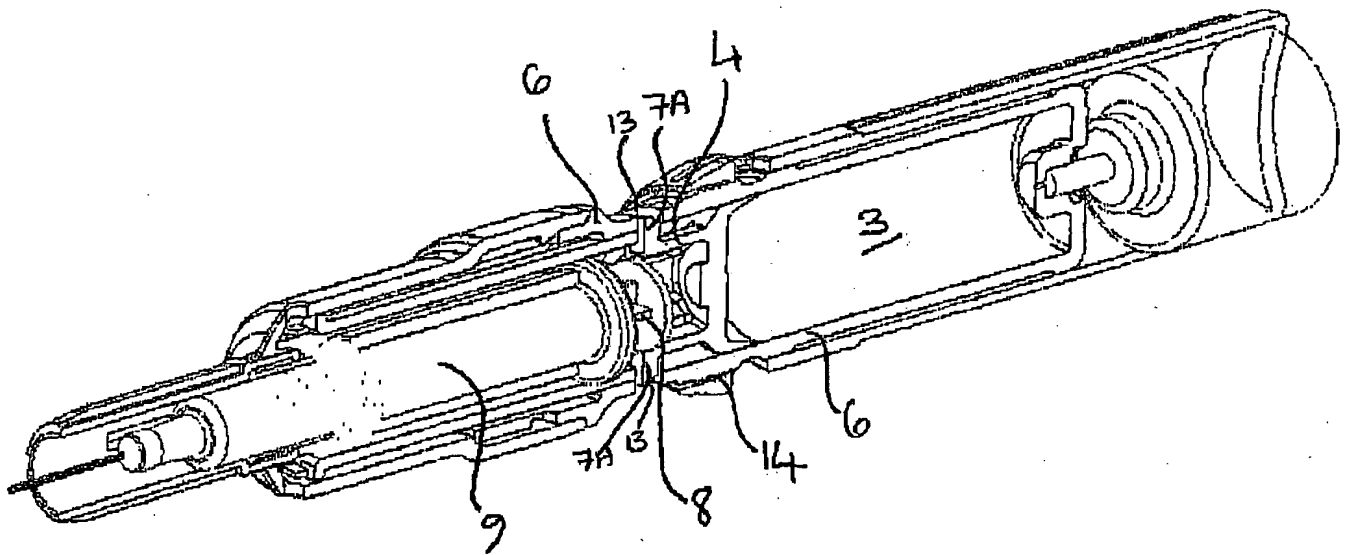


FIGURE 5

6/12

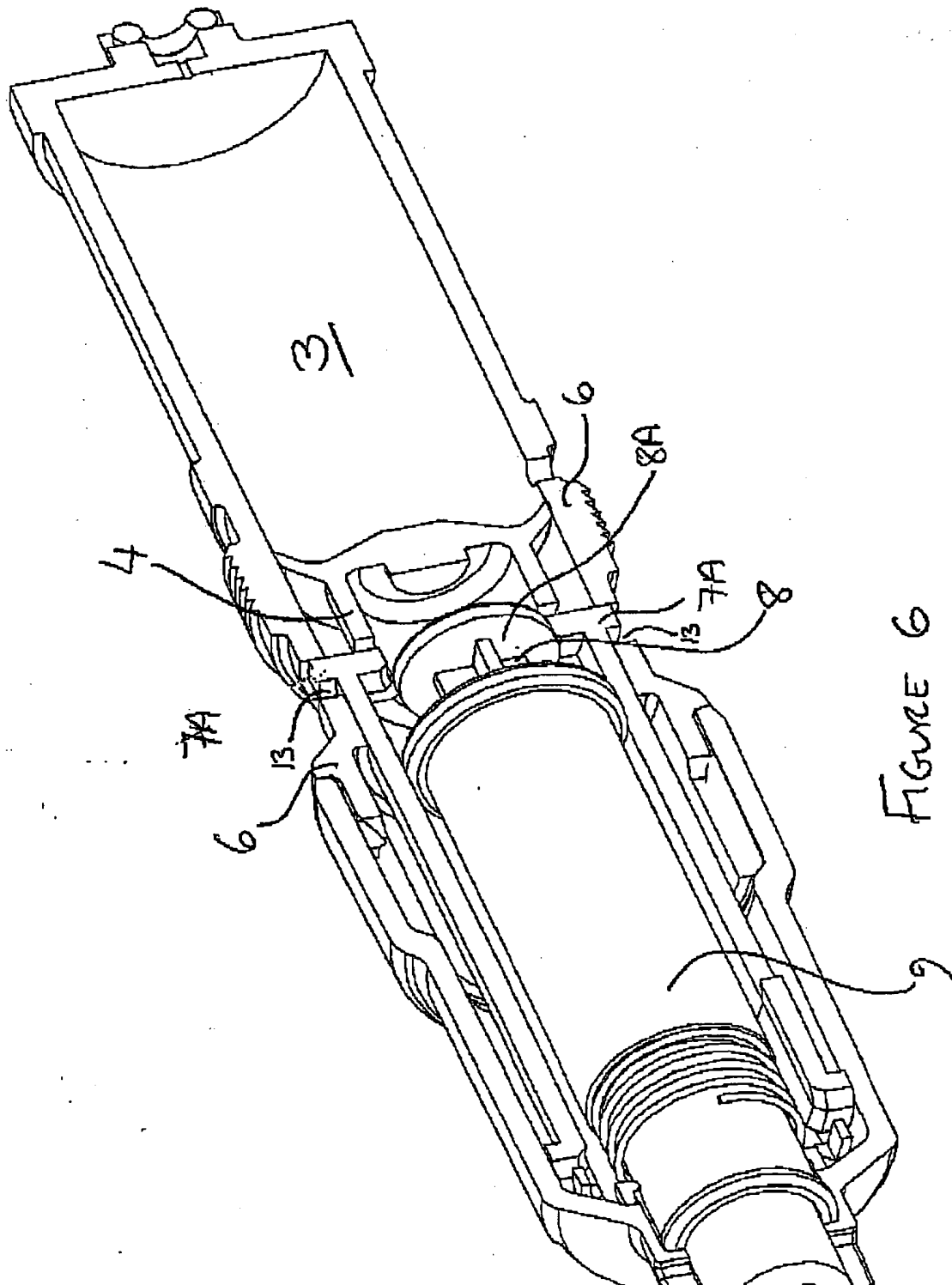


FIGURE 6

7/12

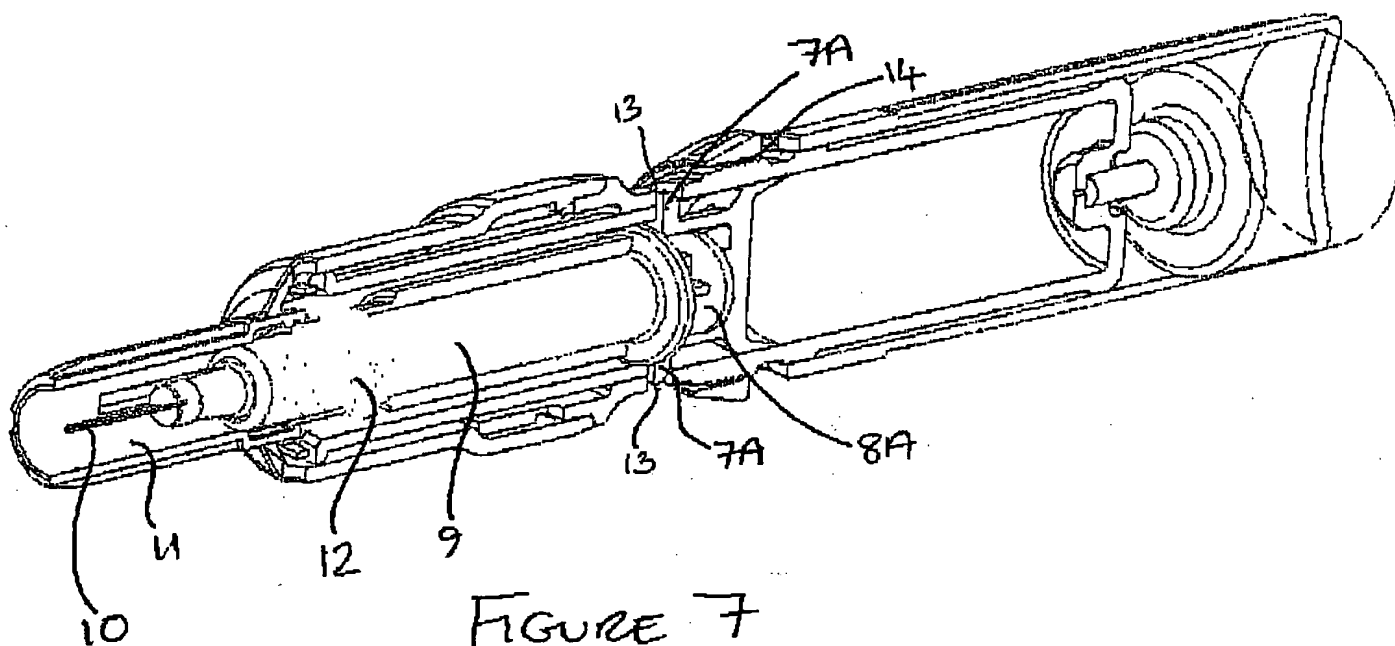


FIGURE 7

8/12

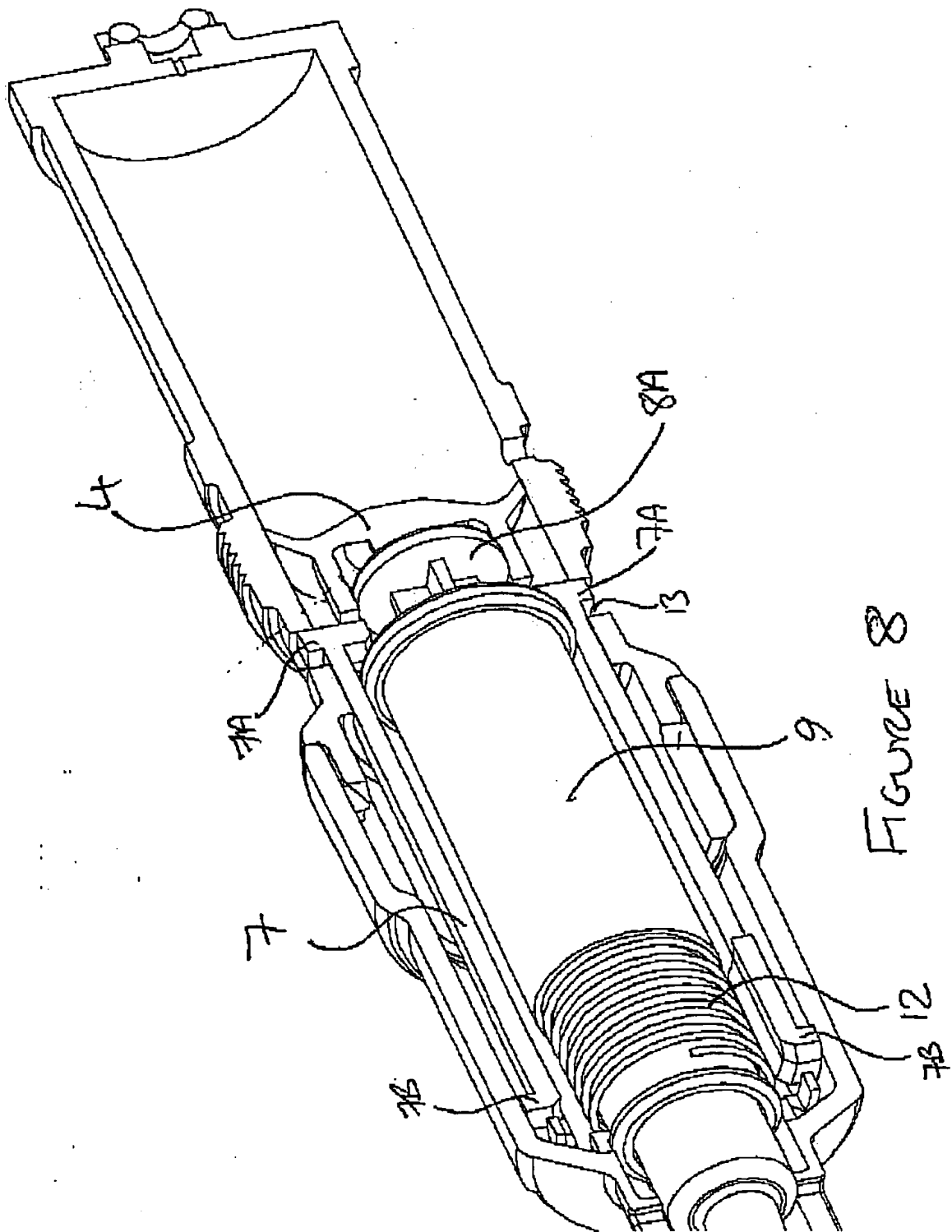


FIGURE 8

RECEIVED 2004-01-23

9/12

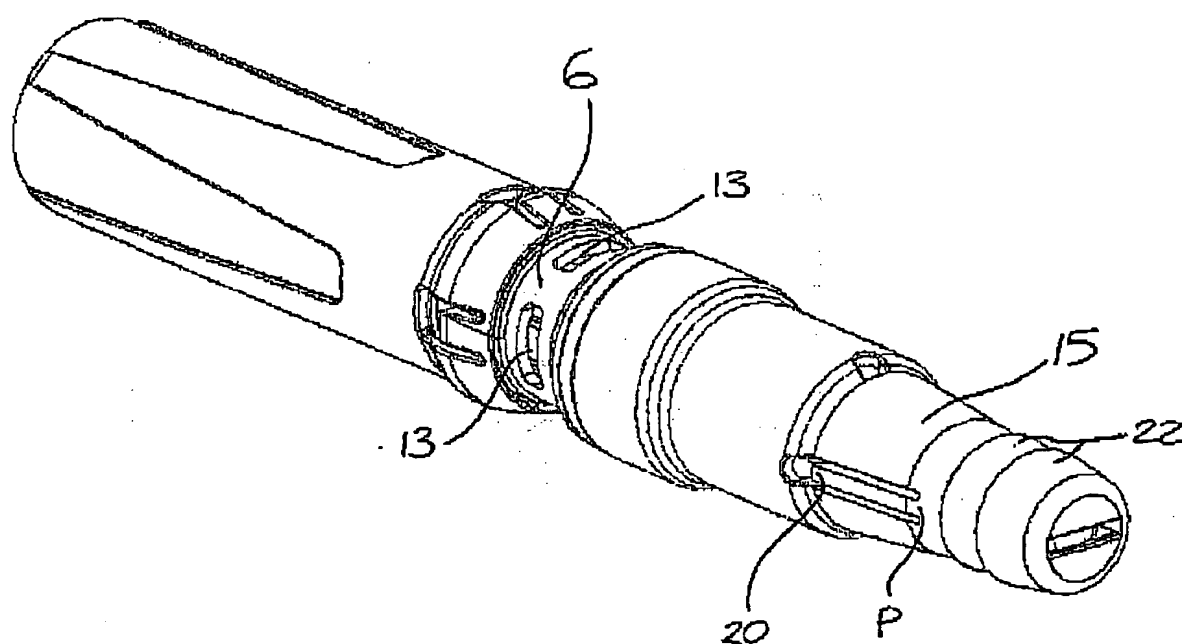


FIGURE 9

10/12

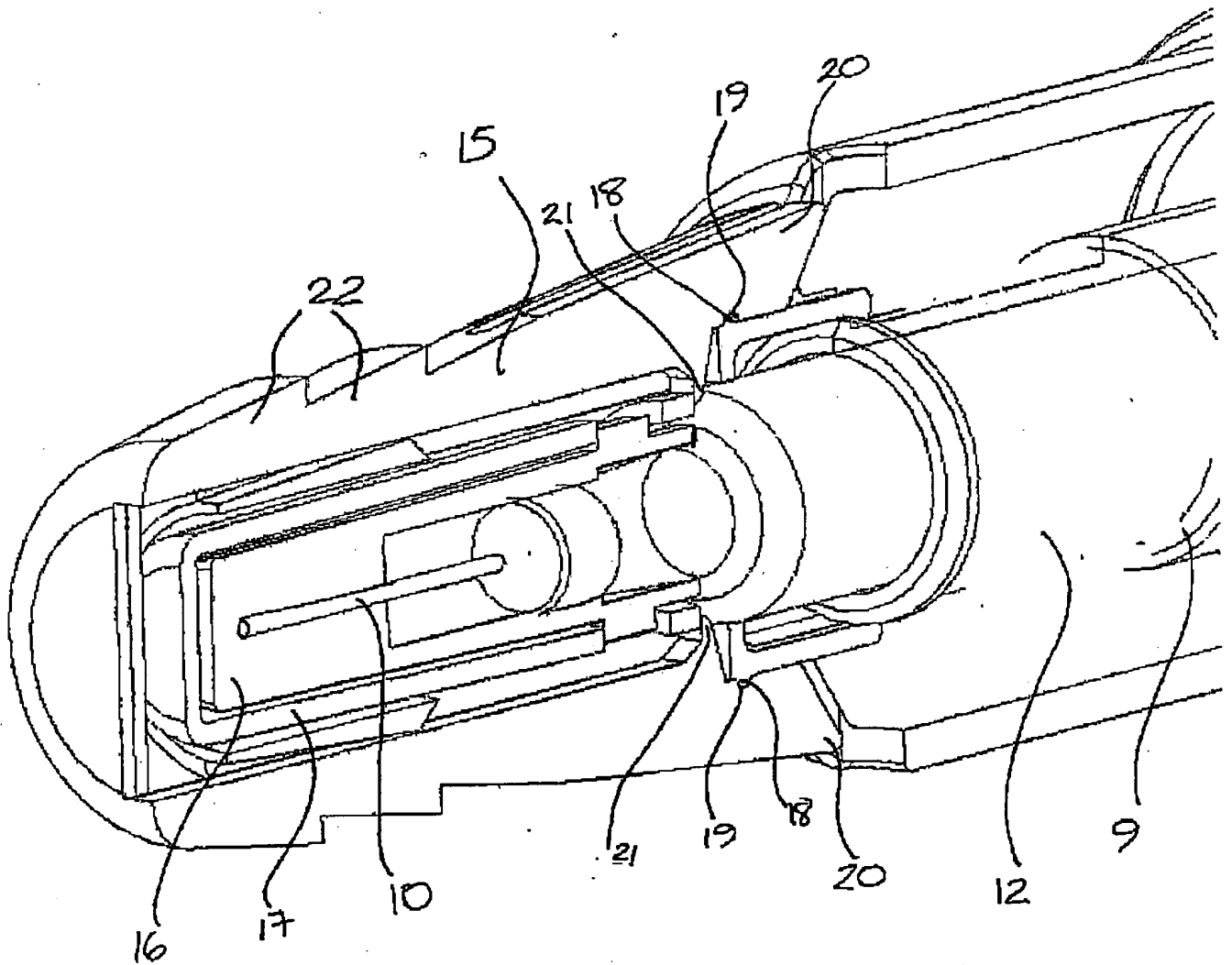


FIGURE 10

11/12

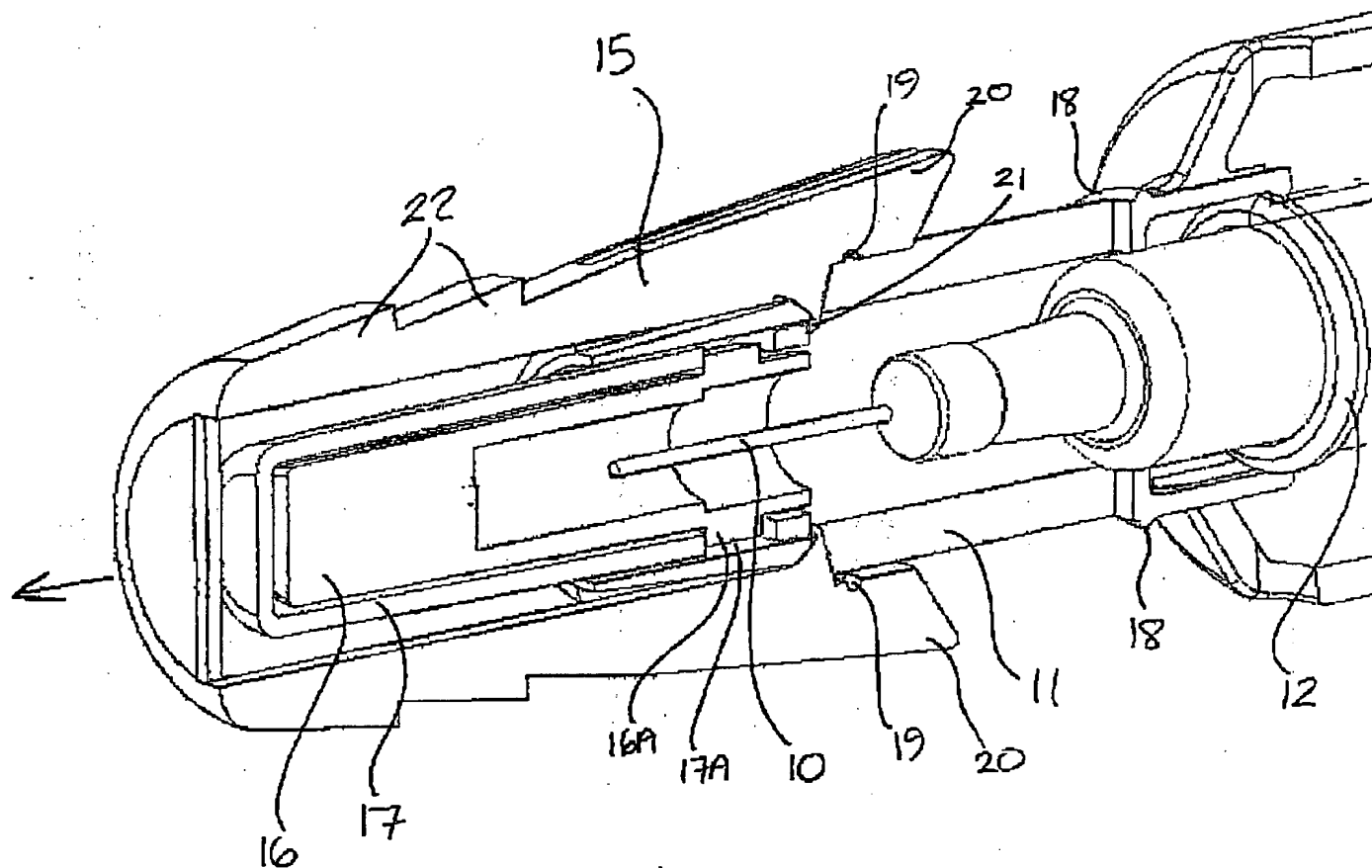


FIGURE 11



12/12

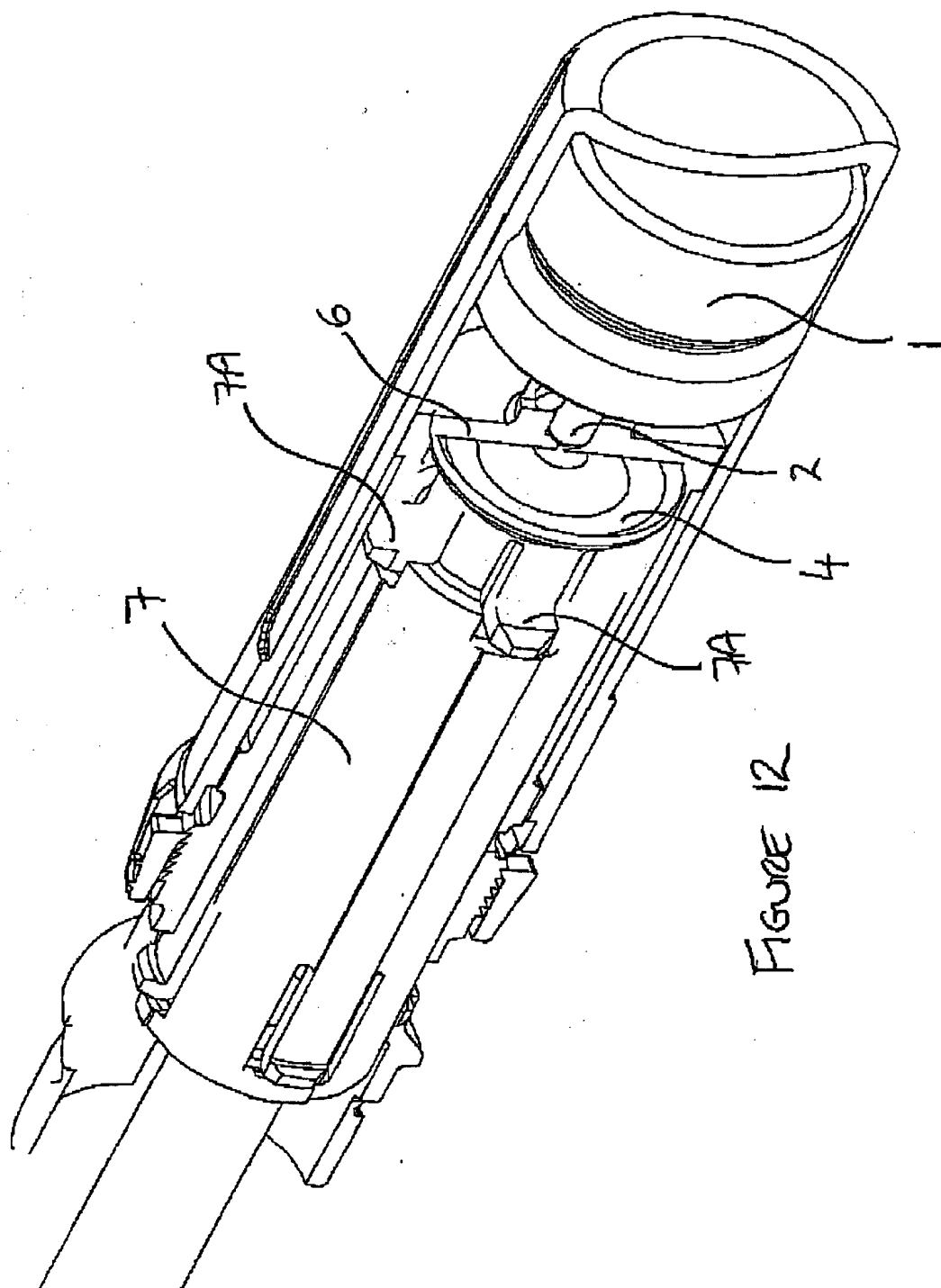


FIGURE 12